# DG BODY ANTIBACTERIAL HAND- triclos an soap DOLGENCORP INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **DRUG FACTS**

## **Active ingredient**

Triclosan 0.15%

## Purpose

Antibacterial

#### Uses

Helps reduce bacteria on the skin

### Warnings

For external use only

## When using this product

avoid contact with eyes. If contact occurs, rinse with water

Stop using this product and ask a doctor if

irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

#### **Directions**

Apply onto wet hands, work into a lather. Rinse thoroughly

#### Other information

Store at room temperature

## **Inactive ingredients**

Water (Aqua), Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Cocamidopropyl Hydroxysultaine, Glycerin, Fragrance (Parfum), Polyquaternium-7, PPG-2 Hydroxyethyl Coco/Isostearamide, DMDM Hydantoin, Tetrasodium EDTA, Citric Acid, Yellow 5 (CI 19140), Red 4 (CI 14700)

### **Label Copy**



## **Drug Facts** Active ingredient Purpose Triclosan 0.15%... Antibacterial Uses - Helps reduce bacteria on the skin. Warnings For external use only When using this product a avoid contact with eyes. If contact occurs, rinse with water. Stop using this product and ask a doctor if irritation or redness develops and lasts. Keep out of reach of children. In case of accidental ingestion, get medical

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"This product is not manufactured or distributed by The Dial Corporation. a Henkel Company, owner of the registered trademark Dial\*.

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DISTRIBUTED BY DOLGENCORP, LLC 100 MISSION RIDGE, GOODLETTSVILLE, TN 37072 MADE IN CANADA

## DG BODY ANTIBACTERIAL HAND

triclosan soap

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55910-294

**Route of Administration** TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength	
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	1.5 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390 UAPO)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDO PRO PYL HYDRO XYSULTAINE (UNII: 62V75NI93W)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
PPG-2 HYDROXYETHYL COCO/ISOSTEARAMIDE (UNII: EK4J71ZKEQ)	
<b>DMDM HYDANTO IN</b> (UNII: BYR0546 TOW)	
EDETATE SO DIUM (UNII: MP1J8420 LU)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:55910-294- 07	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/26/2015		

# Labeler - DOLGENCORP INC (068331990)

# Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture (559 10 - 294)	

Revised: 8/2015 DOLGENCORP INC